IN THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application.

Complete Listing of Claims:

- 1. (Currently amended) A transposon TnKGloxP, characterized in comprising the outer end transposase recognition sequence[[s]] having a base sequence of SEQ ID NO: 3 on one end of the transposon, [[its]] the reverse-complementary sequence of SEQ ID NO: 3 on the other end of the transposon, the loxP site expressed as sequence of SEQ ID NO: 4, the kanamycin resistance (Km^R) gene expressed as sequence of SEQ ID NO: 5 and green fluorescent protein (GFP) gene expressed as sequence of SEQ ID NO: 6.
- 2. (Currently amended) The transposon TnKGloxP according to claim 1, characterized in comprising the base sequence of SEQ ID NO: 1.
 - 3. (Canceled)
- 4. (Currently amended) [[The]] A transposon TnCloxP according to claim 3, characterized in comprising the base sequence of SEQ ID NO: 2.
- 5. (Currently amended) A method for constructing novel strains containing deletion of a specific chromosomal site, characterized in comprising the steps of:
 - (a) preparing two transposons comprising outer end transposase recognition sequences, loxP site and different selectable markers a first transposon according to claim 1 or claim 2 and a second transposon according to claim 4,
 - (b) inserting the above two transposons said first and second transposons,
 respectively, into random positions of different microbial chromosomes
 and determining the each inserted sites;
 - (c) integrating the two microbial chromosomes by P1 phage transduction to position the two transposons comprising different selectable markers first and second transposons on one chromosome; and
 - (d) deleting a chromosomal site between the two loxP sites by expressing Cre gene through from a Cre expression vector introduced.

- 6. (Currently amended) The method for constructing novel strains according to claim 5, wherein the above two transposons are transposon TnKGloxP first transposon is a transposon according to claim 2 1 or 2, or transposon TnCloxP according to claim 3 or 4.
 - 7. (Canceled)
- 8. (Currently amended) The method for constructing novel strains according to claim 5 or 7, characterized in additionally comprising the steps of:

-selecting two mutants from the mutants containing deletion of a specific chromosomal site, and performing P1 phage transduction using one of the selected mutants as the donor and the other as recipient, to constructing construct a new mutant containing all chromosomal deletion sites of the above two mutants;

-using the above obtained mutant again as P1 phage recipient, and the already prepared mutant containing deletion of a specific chromosomal site as donor to perform P1 phage transduction continuously and repeatedly; and

-removing the chromosomal deletion site of other donor mutant from the chromosome of the obtained mutant continuously to reduce the chromosome of the obtained mutant by degrees.